WHAT IS CLAIMED IS:

| 1. An isolated polypeptide comprisin | 1. | An isolate | d polypeptide | comprising |
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- a) the amino acid sequence of SEQ ID NO: 8;
- b) the amino acid sequence of SEQ ID NO: 6, or
- c) the amino acid sequence of SEQ ID NO: 2.
- 2. An antigenic polypeptide comprising:
- a) an immunogenic amino acid sequence exhibiting

 10 identity overall length of at least 12 amino acids to SEQ

 ID NO: 8;
 - b) an immunogenic amino acid sequence exhibiting identity over a length of at least 12 amino acids to SEQ ID NO: 6; or
- 15 c) an immunogenic amino acid sequence exhibiting identity over a length of at least 12 amino acids to SEQ ID NO: 2.
 - 3. An antigenic polypeptide of:
- 20 a) Claim 2a, further comprising:
 - i) a second Tength of identity of 12 amino acids;
 - ii) a detection or purification tag;
 - iii) a sequence of another chemokine receptor;

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- iv) a carbohydrate;
- b) Claim 2b, further comprising:
 - i) a second length of identity of 12 amino
 acids;
- ii) a detection or purification tag;
 - iii) a sequence of another chemokine receptor;

or

- iv) a carbohydrate; or
- c) Claim 2c, further comprising:
- i) a second length of identity of 12 amino acids;
 - ii) a detection or purification tag;
 - iii) a sequence of another chemokine; or

- 4. The polypeptide of Claim 1, which;
- a) has a molecular weight of at least 3 kD with
- 5 natural glycosylation;
 - b) is a synthetic polypeptide;
 - c) is attached to a solid substrate;
 - d) is conjugated to another chemical moiety;
 - e) is a 5-fold or less substitution from natural
- 10 sequence; or
 - f) is a deletion or insertion variant from a natural sequence.
 - 5. A composition comprising:
- a) a sterile\polypeptide of Claim 1a,
 - b) a sterile polypeptide of Claim 1b; or
 - c) a sterile palypeptide of Claim 1c.
 - 6. A kit comprising a polypeptide of Claim 1, and:
 - a) a compartment comprising said polypeptide; and/or
 - b) instructions for use or disposal of reagents in said kit.
 - 7. A method of using said polypeptide of Claim 1
- 25 to:

- a) produce an antiserum, comprising immunizing an animal with said polypeptide, and isolating said antiserum; or
- b) produce an antibody:antigen complex, comprising 30 contacting said polypeptide with a specific antibody, thereby producing said complex.
 - 8. A binding compound comprising an antigen binding portion from an antibody, which specifically binds to a polypeptide of Claim 1, wherein:
 - a) said binding compound is an Fv, Fab, or Fab2 fragment;

- b) said binding compound is conjugated to another chemical moiety; or
 - c) said antibody:
 - i) is raised against a peptide sequence of a mature polypeptide of Figure 1 or Figures 3A-3C;
 - ii) is raised against a peptide sequence of a
 mature rodent polypeptide of Figure 5;
 - iii) is immunoselected;
 - iv) is a polyclonal antibody;
- v) binds to a denatured rodent CXC N4, rodent DNAXCCR10, or primate BLRx;
 - vi) exhibits a Kd to antigen of at least 30 μ M;
 - vii) is attached to a solid substrate, including a bead or plastic membrane;
 - viii) is/in a sterile composition; or
 - ix) is detectably labeled, including a radioactive or fluorescent label.
- 9. A kit comprising said binding compound of Claim 20 8, and:
 - a) a compartment comprising said binding compound;
 and/or
 - b) instructions for use or disposal of reagents in said kit.
- 10. A composition comprising:
 - a) a sterile binding compound of Claim 8; or
 - b) said binding compound of Claim 8 and a carrier, wherein said carrier is:
 - i) an aqueous compound, including water, saline, and/or buffer; and/or
 - ii) formulated for oral, rectal, nasal, topical, or parenteral administration.
 - 11. An isolated or recombinant nucleic acid encoding a polypeptide of Claim 1, wherein said nucleic acid:

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a) encodes an antigenic peptide sequence of Figure 1or Figures 3A-3C;

- b) encodes an antigenic rodent peptide sequence of Figure 5;
- c) encodes a plurality of antigenic peptide sequences of Figure or Figures 3A-3C;
- d) encodes a plurality of antigenic peptide sequences of Figures 2A-2B;
- e) exhibits identity of at least 27 nucleotides of SEQ ID NO: 7, 5, or 1;
 - f) is an expression vector;
 - g) further comprises an origin of replication;
 - h) is from a natural source;
 - i) comprises a detectable label;
- j) comprises synthetic nucleotide sequence;
 - k) is Yess than 6 kb, preferably less than 3 kb;
 - 1) is from a mammal, including a rodent;
 - m) /comprises a natural full length coding sequence;
 - n) is a hybridization probe for a gene encoding said
- 20 protéin; or
 - o) is a PCR primer, PCR product, or mutagenesis primer.
- 12. A cell or tissue comprising a recombinant 25 nucleic acid of Claim 11.
 - 13. The cell of Claim 12, wherein said cell is:
 - a) a prokaryotic cell;
 - b) a eukaryotic cell;
- 30 c) a bacterial cell;
 - d) a yeast cell;
 - e) an insect cell;
 - f) a mammalian cell;
 - g) a mouse cell;
- 35 h) a primate cell; or
 - i) a human cell.

- 14. A kit comprising said nucleic acid of Claim 11, and:
 - a) a compartment comprising said nucleic acid;
- b) \ a compartment further comprising a polypeptide of SEQ ID NO: 8, 6, or 2; and/or
- c) instructions for use or disposal of reagents in said kit.
 - 15. A nucleic acid which:
- a) hybridizes under wash conditions of 45° C and less than 700 mM salt to SEQ ID NO: 1;
- b) hybridizes under wash conditions of 45° C and less than 700 mM salt to SEQ ID NO: 5;
- c) hybridizes under wash conditions of 45° C and 15 less than 700 mM salt to SEQ ID NO: 7;
 - d) exhibits identity over a stretch of 30 nucleotides to SEQ ID NO: 7;
 - e) exhibits identity over at least 30 nucleotides to SEQ ID NO: 5; or $\fill \fill \fill$
- 20 f) exhibits identity over at least 30 nucleotides to SEQ ID NO 1.
 - 16. The nucleic acid of Claim 15, wherein:
- a) said wash conditions are at 55° C and/or 500 mM 25 salt; or
 - b) said identity is over at least 55 nucleotides.
 - 17. $\dot{}$ The nucleic acid of Claim 16, wherein:
- \sim 50 mM a) said wash conditions are at 65° C and/or 150 mM \sim 30 salt; or \sim
 - b) said identity is over at least 75 nucleotides.
 - 18. A kit comprising said nucleic acid of Claim 15, and:
 - a) a compartment comprising said nucleic acid;
 - b) a compartment further comprising a polypeptide of SEQ ID NO: 8, 6, or 2; and/or

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instructions for use or disposal of reagents in said kit.

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- 19. A method of using said nucleic acid of Claim 15:
- a) to produce a duplex nucleic acid, comprising contacting one strand of the nucleic acid to the complementary strand, thereby producing said duplex; or
- b) to produce a polypeptide, comprising expressing said nucleia acid in a host cell, thereby producing said 10 polypeptide.
 - \20. A method of screening for a compound which binds to a polypeptide of Claim 1 having SEQ ID NO: 8, comprising contacting said compound to said polypeptide, and detecting binding.
 - An isolated polypeptide, comprising the amino acid sequence of SEQ ID NO:8, or a polypeptide having at least about 80% sequence homology thereto.
 - An isolated polynucleotide encoding the polypeptide of claim 21.
 - The polynucleotide of claim 22, wherein the polynucleotide comprises the nucleotide sequence of SEQ ID NO:7, or a polynucleotide having at least about 80% sequence homology thereto.
 - 24. A recombinant vector comprising
 - (a) a polynucleotide according to claim 22; and
 - (b) control elements that are operably linked to said polynucleotide whereby a coding sequence within said polynucleotide can be transcribed and translated in a host cell, and at least one of said control elements is heterologous to said coding sequence.

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- 25. A host cell transformed with the recombinant vector of claim 24.
- 26. A method of producing a recombinant polypeptide comprising:

- (a) providing a population of host cells according to claim 25; and
- (b) culturing said population of cells under conditions whereby a polypeptide encoded by the coding
 sequence present in said recombinant vector is expressed.
 - 27. A method of expressing a recombinant polypeptide comprising:
- (a) transforming a host cell with the recombinant vector of claim 22; and
 - (b) causing expression of a polypeptide encoded by the coding sequence present in said recombinant vector.
- - 29. The method of claim 28, wherein the host cell is in the region of a wound.
- 25 30. A method of treating a wound comprising:
 - (a) transforming a host cell in vivo with the polynucleotide of claim 22, wherein the host cell is in the region of a wound; and
- (b) causing expression of a polypeptide encoded by the coding sequence present in said recombinant vector.
 - 31. A method of treating a wound comprising modulating the *in vivo* expression of an endogenous polynucleotide in the region of the wound, wherein the polynucleotide encodes a polypeptide comprising the amino acid sequence of SEQ ID NO:8.

- 32. The method of claim 31, wherein expression is up-regulated.
- 33. An antibody reactive with the polypeptide of 5 claim 21.
 - 34. The antibody of claim 33, wherein the antibody is a polyclonal antibody.
- 10 35. The antibody of claim 33, wherein the antibody is a monoclonal antibody.
- 36. A method of treating a wound comprising administering the antibody of claim 33 to a subject in need thereof.

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